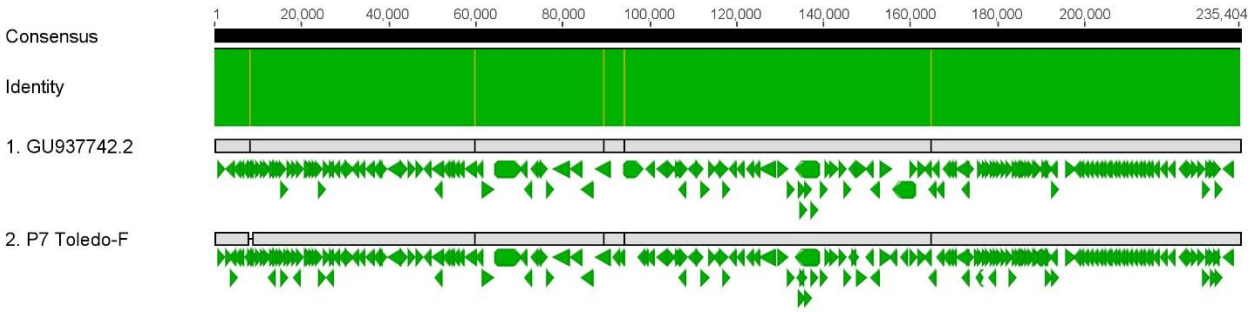
A



B

Sequence aligned to	Coordinate	Type
GU937742.2	8274	insertion of G in non-coding region (likely homopolymer)
GU937742.2	59592	SNP A to C causes L to W in UL45
GU937742.2	89517	SNP G to C synonymous mutation G to G
GU937742.2	94188	SNP T to C synonymous mutation P to P
GU937742.2	94227	SNP T to G synonymous mutation A to A
GU937742.2	164390	SNP A to G synonymous mutation S to S

Figure S1